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APPLICATION NO.	F	ILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/426,792		10/22/1999	DENNIS T. MANGANO	9114-004-999	2354
20583	7590	01/29/2004	•	EXAMINER	
JONES DA		CET	SPIVACK, PHYLLIS G		
222 EAST 41ST STREET NEW YORK, NY 10017				ART UNIT	PAPER NUMBER
				1614	
				DATE MAILED: 01/29/2004	1

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)					
Office Action Summary	09/426,792	MANGANO, DENNIS T.					
Office Action Summary	Examiner	Art Unit					
The MAILING DATE of this communication on	Phyllis G. Spivack	1614					
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with th	e correspondence address					
A SHORTENED STATUTORY PERIOD FOR REPL THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.1 after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a repl - If NO period for reply is specified above, the maximum statutory period - Failure to reply within the set or extended period for reply will, by statute - Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b). Status	36(a). In no event, however, may a reply be y within the statutory minimum of thirty (30) will apply and will expire SIX (6) MONTHS fr a, cause the application to become ABANDO	e timely filed days will be considered timely. om the mailing date of this communication. NED (35 U.S.C. § 133).					
1) Responsive to communication(s) filed on 23 C	<u> 0ctober 2003</u> .						
2a) This action is FINAL . 2b) ⊠ This	action is non-final.						
	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.						
Disposition of Claims							
4) Claim(s) 1-6,13-16 and 49-54 is/are pending in 4a) Of the above claim(s) is/are withdray 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-6, 13-16, 49-54 are subject to restrict to the strict of the str	wn from consideration.	ent.					
Application Papers							
9) The specification is objected to by the Examiner.							
10) The drawing(s) filed on is/are: a) acc							
Applicant may not request that any objection to the							
Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex							
Priority under 35 U.S.C. §§ 119 and 120	Administration and all additions of the	007.00.01.01.00.1.01.01.01.01.01.01.01.01.0					
12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).							
a) All b) Some * c) None of: 1. Certified copies of the priority document 2. Certified copies of the priority document 3. Copies of the certified copies of the prio application from the International Burea * See the attached detailed Office action for a list 13) Acknowledgment is made of a claim for domest since a specific reference was included in the fir 37 CFR 1.78. a) The translation of the foreign language pro 14) Acknowledgment is made of a claim for domest reference was included in the first sentence of the	ts have been received. Its have been received in Application of the certified copies not receive priority under 35 U.S.C. § 11 st sentence of the specification ovisional application has been ric priority under 35 U.S.C. §§ 1	ation No ived in this National Stage ived. 9(e) (to a provisional application) or in an Application Data Sheet. received. 20 and/or 121 since a specific					
Attachment(s)							
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449) Paper No(s) _ 	5) Notice of Informa	ary (PTO-413) Paper No(s) al Patent Application (PTO-152)					

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RESTRICTION

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Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is a β_1 -adrenergic selective blocking agent.
- II. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is an α_2 agonist.
- III. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is a nitrate.
- IV. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is a calcium channel blocker.
- V. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is an ACE inhibitor.

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VI. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is a platelet inhibitor.

VII. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is a thrombosis inhibitor.

VIII. Methods for reducing cardiovascular disease complications following surgery under defined conditions, comprising administering a pharmacologic cardiovascular agent, wherein the cardiovascular agent is an agent not encompassed in one of the Groups *supra*.

The inventions are distinct, each from the other, for the following reasons:

The Groups have acquired a separate status in the art. Depending on the particular agent contemplated, the Groups would be separately classified. A search for methods of use, comprising the administration of cardiovascular agents such as nitrates, does not suggest methods of use wherein an α_2 agonist is administered. The searches are not co-extensive. The Groups encompass a plethora of compounds resulting in an unreasonable search burden. Thus restriction for examination purposes, as indicated, is proper.

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Applicant is advised that to be complete, the reply to this requirement must include an election of the invention to be examined even though the requirement is traversed (37 CFR 1.143).

Any inquiry concerning this communication should be directed to Phyllis G.

Spivack at telephone number 703-308-4703.

Phyllis G. Spivack Primary Examiner Art Unit 1614

January 27, 2004

PHYLLIS SPIVACK PRIMARY EXAMINER